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(21) International Application Number: PCT/US89/03341 (22) International Filing Date: 2 August 1989 (02.08.89) (30) Priority data: 227,371 2 August 1988 (02.08.88) US (71) Applicant: BONE CARE INTERNATIONAL, INC. [US/US]; 313 W. Beltline Highway, Madison, WI 53713 (US). (72) Inventors: DeLUCA, Hector, F. ; 1809 Hwy. BB, Deerfield, WI 53531 (US). BISHOP, Charles, W. ; 313 W. Beltline Highway, Madison, WI 53713 (US). MAZESS, Richard, B. ; 2526 Gregory St., Madison, WI 53711 (US). GALLAGHER, John, C. ; 601 North 30th Street, Omaha, NE 68131 (US).		(74) Agent: GULBRANDSEN, Carl, E.; Stroud, Stroud, Wilink, Thompson & Howard, 25 West Main Street, Box 2236, Madison, WI 53701-2236 (US). (81) Designated States: AT (European patent), AU, BE (European patent), CH (European patent), DE, DE (European patent), FR (European patent), GB, GB (European patent), IT (European patent), JP, LU (European patent), NL, NL (European patent), SE, SE (European patent). Published <i>Without international search report and to be republished upon receipt of that report.</i>
(54) Title: METHOD FOR TREATING AND PREVENTING LOSS OF BONE MASS (57) Abstract The invention describes and claims the use of 1 α -Vitamin D2 as a medicament in the treatment of osteoporosis. Active Vitamin D3 compounds are ineffective or minimally effective at dosage ranges which can be tolerated by the human. At dosage ranges which are required to effectively restore lost bone mass, active Vitamin D3 compounds present a serious amount of toxicity in the form of hypercalcemia and hypercalciuria. The inventors have found that 1 α -Vitamin D2 which is equally as active as 1 α -Vitamin D3 in stimulating intestinal calcium absorption can be safely administered at dosages above 3.0 μ g/day without stimulating bone reabsorption.		

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METHOD FOR TREATING AND
PREVENTING LOSS OF BONE MASS

Technical Field

This invention relates to a method for treating and preventing metabolic bone disorders characterized by loss of bone mass or by disproportionately low bone mineral content.

More specifically, this invention relates to a method for treating or preventing various known forms of osteoporosis, e.g. postmenopausal, senile and steroid-induced osteoporosis or other disease states one of the characteristics of which is the loss of bone mass or decreased mineral content.

Still more specifically, this invention relates to a method for treating or for preventing the depletion of bone of women who are entering menopause or who are postmenopausal.

Background of the Invention

Numerous metabolic bone disorders are known to the medical community which are characterized by loss of bone mass or disproportionate mineralization of bone. These disorders include postmenopausal osteoporosis, senile osteoporosis, renal osteodystrophy, corticosteroid-induced osteopenia, and anticonvulsant osteomalacia. Of these disorders, postmenopausal and senile osteoporosis are most commonly encountered in normal medical practice.

As a group, these bone depletive disorders are a major and growing public health problem in the United States. Together, they

cause more than 1 million bone fractures per year, primarily of the spine, hip, and distal forearm, and result in an annual cost of \$6 to 7 billion to the American society. Unfortunately, the incidence of these bone disorders will rise significantly as the mean age of the U.S. population continues to increase.

Despite differing etiologies, the aforementioned metabolic bone disorders develop during an extended period of negative calcium balance. This commonality of the disorders suggests that agents which stimulate intestinal calcium absorption may be effective in restoring calcium balance and thereby treating or preventing the development of such bone disorders. It has long been known that Vitamin D plays a critical role in stimulating calcium absorption and regulating calcium metabolism. The discovery of the active forms of Vitamin D, [M.F. Holick et al., Proc. Natl. Acad. Sci. USA 68, 803-804 (1971); G. Jones et al., Biochemistry 14, 1250-1256 (1975)] and active Vitamin D analogues [M.F. Holick et al., Science 180, 190,191 (1973); H.Y. Lam et al., Science 186, 1038-1040 (1974)], caused much excitement and speculation about the usefulness of these compounds in the treatment of bone depletive disorders.

Animal studies examining the effects of these active Vitamin D compounds suggested that such agents would be useful in restoring calcium balance. Further, an early clinical study indicated that administration of 0.5 μ g/day of 1,25-dihydroxycholecalciferol (1,25 Vitamin D₃) to a group of postmenopausal women improved the intestinal calcium absorption as well as the calcium balance of these women. On this basis, U.S. Patent 4,225,596 (" '596 Patent") described and claimed the use of 1,25 Vitamin D₃ for increasing calcium absorption and retention. Such use also was claimed in the same patent for 1,25 dihydroxyergocalciferol (1,25 Vitamin D₂) and 1 α -hydroxyergocalciferol (1 α -Vitamin D₂), which the patent teaches are "eminently suitable and readily substitutable for the 1,25 dihydroxycholecalciferol."

The best indicator of the efficacy of Vitamin D compounds to prevent or treat depletive bone disorders is bone itself rather than calcium absorption or calcium balance. More recent clinical

data indicates that at the dosage ranges taught in the '596 Patent, 1,25 Vitamin D₃ has, at best, modest efficacy in preventing or restoring loss of bone mass or bone mineral content [S.M. Ott, C.H. Chesnut, In: J Jensen et al, ed, Norhaven A/S, Viborg, p.83 (1987); J.C. Gallagher et al., 7th Workshop on Vitamin D, Rancho Mirage, p. 196 (1988); J. Aloia et al., Amer. J. Med. 84:401-08 (1988)].

Together these clinical studies with 1,25 Vitamin D₃, and one other conducted with 1 α -Vitamin D₃ [M. Shiraki et al., Endocrinol. Japan 32, 305-315 (1985)], indicate that the ability of these agents to restore lost bone mass or bone mineral content is dose related. These studies also indicate, however, that at the dosage ranges required for the agents to be truly effective, toxicity in the form of hypercalcemia and hypercalciuria becomes a major problem. Thus, attempts to increase the amount of 1,25 Vitamin D₃ above 0.5 μ g/day have frequently resulted in toxicity. At dosage levels below 0.5 μ g/day no effects are observed on bone. [See G.F. Jensen et al., Clin. Endocrinol. 16, 515-524 (1982); C. Christiansen et al., Eur. J. Clin. Invest. 11, 305-309 (1981)]. Two μ g/day of 1 α -Vitamin D₃ was found to have efficacy in increasing bone mass in patients exhibiting senile osteoporosis [O.H. Sorensen et al., Clin. Endocrinol. 7, 169S-175S (1977)]. Data from the clinical studies in Japan, a population that has low calcium intake, indicate that efficacy is found with 1 α -Vitamin D₃ when administered at 1 μ g/day [M. Shiraki et al., Endocrinol. Japan. 32:305-315 (1985); H. Orimo et al., Bone and Mineral 3, 47-52 (1987)]. At 2 μ g/day, however, toxicity with 1 α -Vitamin D₃ occurs in approximately 67 percent of the patients, and at 1 μ g/day this percentage is approximately 20 percent.

Thus, the prior art teaches that due to their toxicity, 1 hydroxylated Vitamin D compounds can only be administered at dosages that are, at best, modestly beneficial in preventing or treating loss of bone or bone mineral content. Indeed, Aloia recommends that alternative routes of administration be sought which might avoid the toxicity problems and allow higher dosage levels to be achieved. [J.Aloia et al, Amer J Med 84:401-408 (1988)].

Summary of the Invention

The present invention discloses a method whereby the bone forming activity of 1-hydroxylated Vitamin D compounds is made available to safely prevent or treat bone depletive disorders. With the exception of the data provided herein, no clinical data are available with respect to the 1-hydroxylated Vitamin D₂ compound. Nonetheless, in the prior art, such compounds have been considered as essentially equivalent to their Vitamin D₃ counterparts. Thus, when the prior art teaches that because of its extreme toxicity 1 α -Vitamin D₃ may not be safely administered at normal calcium intakes in excess of 2 μ g/day, a person skilled in the art would understand that the same warning also applies to 1 α -Vitamin D₂.

During the course of our investigations, we have compared 1 α -Vitamin D₂ to 1 α -Vitamin D₃. 1 α -Vitamin D₂ is equally active as 1 α -Vitamin D₃ in the healing of rickets, the stimulation of intestinal calcium absorption and in the elevation of serum inorganic phosphorous of rachitic rats. [G. Sjoden et al., J. Nutr. 114, 2043-2946 (1984)]. In the same laboratory animal we also have found that, 1 α -Vitamin D₂ is 5 to 15 times less toxic than 1 α -Vitamin D₃. [G. Sjoden et al., Proc. Soc. Exp. Biol. Med. 178, 432-436 (1985)].

It has now been found that 1 α -Vitamin D₂ may be safely administered to human subjects experiencing or having a tendency toward loss of bone mass or bone mineral content at dosages greater than 3 μ g/day.

The present invention is intended to be used in all bone depletive disorders which respond to administration of active forms of Vitamin D, including renal osteodystrophy, anticonvulsant osteomalacia, hypophosphatemic Vitamin D-resistant rickets, pseudo-deficiency (Vitamin D-dependent) rickets, nutritional and malabsorptive rickets, osteomalacia and osteopenias secondary to hypoparathyroidism post-surgical hypoparathyroidism, idiopathic hypoparathyroidism, pseudo-parathyroidism and alcoholism.

The present invention also is intended to be used in those bone depletive disorders classified as osteoporosis, particularly post-menopausal osteoporosis, senile osteoporosis, idiopathic osteoporosis, immobilization osteoporosis, post-lactational osteoporosis, glucocorticoid, alcohol or drug-induced osteoporosis, juvenile osteoporosis, osteoporosis secondary to gonadal insufficiency, malnutrition, hyperprolactinemia, or disorders of the gastrointestinal tract, liver, or kidneys, and osteoporosis that is a sequela of prior osteomalacia, chronic disorders involving the bone marrow, and heritable forms of osteoporosis such as osteoporosis imperfecta and its variants, and other heritable disorders of connective tissue.

Example 1

Six postmenopausal osteoporotic women were enrolled in an open label study. The selected patients had ages between 55 and 75 years, and exhibited L2-L3 vertebral bone mineral density between 0.7 and 1.05 g/cm², as determined by measurements with a LUNAR Radiation dual photon absorptiometer. (The mean bone mineral density in women with osteoporosis is about 0.85 ± 0.17 g/cm², so that these limits correspond to about the 15th to the 85th percentiles.)

On admission to the study, all patients received instruction on selecting a daily diet containing 400 to 600 mg. of calcium. Compliance to this diet was verified at weekly intervals by 24-hour food records and by interviews with each patient.

All patients completed a one-week baseline period, a five-week treatment period, and a one-week post-treatment observation period. During the treatment period, patients orally self-administered a 1 α -Vitamin D₂ at an initial dose of 0.5 μ g/day for the first week, and at successively higher doses of 1.0, 2.0, 4.0 and 5.0 μ g/day in each of the following four weeks. All doses were administered before breakfast.

Blood and urine chemistries were monitored on a weekly basis throughout the study. Key blood chemistries included fasting serum levels of calcium, phosphorus, osteocalcin, creatinine blood urea nitrogen. Key urine chemistries included 24-hour excretion of calcium, phosphorus and creatinine.

Data from the study clearly demonstrated that 1α -Vitamin D₂ can be safely administered at high dose levels. As shown in Table 1, all six patients tolerated a 4.0 μ g/day dose of this compound without exhibiting hypercalciuria (>350 mg/24 hr) or hypercalcemia (>11.0 mg/dL). Five of the patients tolerated a 5.0 μ g/day dosage without side effects, while the sixth patient showed a mild hypercalciuria, without attendant hypercalcemia, at this dosage.

TABLE 1

Dose-response relationship between orally administered 1α -Vitamin D₂ and fasting serum and 24-hour urinary calcium values in postmenopausal osteoporotic patients.

Dose	Fasting Serum Ca		24-hr Urinary Ca	
	Mean \pm SE	Range	Mean \pm SE	Range
0.5 μ g/d	9.80 \pm .10	(9.51-10.20)	121.8 \pm 17.2	(55.3-180.0)
1.0 "	9.81 \pm .08	(9.61-10.18)	132.2 \pm 16.7	(67.8-187.1)
2.0 "	9.87 \pm .15	(9.58-10.56)	169.9 \pm 16.4	(110.5-215.1)
4.0 "	9.92 \pm .10	(9.65-10.33)	193.9 \pm 30.1	(133.8-324.1)
5.0 "	9.80 \pm .09	(9.61-10.15)	221.3 \pm 37.0	(149.9-405.9)

Additional blood and urine data from this clinical study supported the conclusion that 1α -Vitamin D₂ can be safely administered at high dosages. In particular, this compound did not adversely affect kidney function, as determined by creatinine clearance and blood levels of urea nitrogen; nor did it increase urinary excretion of hydroxyproline, indicating the absence of any stimulatory effect on bone resorption. The compound had no effect on any routinely monitored serum parameters, indicating the absence of adverse metabolic effects.

A positive effect of 1α -Vitamin D₂ on calcium homeostasis was evident from modest increases in 24-hour urinary calcium levels (see Table 1), confirming that the compound increases intestinal calcium absorption.

Example 2

A clinical study is conducted with postmenopausal osteoporotic out-patients having ages between 55 and 75 years. The study involves up to 120 patients randomly divided into three treatment groups, and continues for 12 to 18 months. Two of the treatment groups receive constant dosages of 1α -Vitamin D₂ (u.i.d.; two different dose levels above 3.0 μ g/day) and the other group receives a matching placebo. All patients maintain a normal intake of dietary calcium (500 to 800 mg/day) and refrain from using calcium supplements. Efficacy is evaluated by pre- and post-treatment comparisons of the patient groups with regard to (a) total body calcium retention, (b) radial and spinal bone mineral density as determined by dual-photon absorptiometry (DPA) or dual-energy x-ray absorptiometry (DEXA), (c) bone biopsies of the iliac crest, and (d) determinations of serum osteocalcin. Safety is evaluated by comparisons of urinary hydroxyproline excretion, serum and urine calcium levels, creatinine clearance, blood urea nitrogen, and other routine determinations.

The results show that patients treated with 1α -Vitamin D₂ exhibit significantly higher total body calcium, and radial and spinal bone densities relative to patients treated with placebo. The treated patients also exhibit significant elevations in serum osteocalcin. Bone biopsies obtained from the treated patients show that 1α -Vitamin D₂ stimulates normal bone formation. The monitored safety parameters confirm an insignificant incidence of hypercalcemia or hypercalciuria, or any other metabolic disturbance with 1α -Vitamin D₂ therapy.

Example 3

A clinical study is conducted with healthy postmenopausal women having ages between 55 and 60 years. The study involves up to 80 patients randomly divided into two treatment groups, and continues for 12 to 18 months. One treatment group receives a constant dosage of 1α -Vitamin D₂ (u.i.d.; a dose level above 3.0 μ g/day) and the other receives a matching placebo. The study is conducted as indicated in Example 2 above.

The results show that patients treated with 1α -Vitamin D₂ exhibit reduced losses in total body calcium, radial or spinal bone densities relative to baseline values. In contrast, patients treated with placebo show significant losses in these parameters relative to baseline values. The monitored safety parameters confirm the safety of long-term 1α -Vitamin D₂ administration at this dose level.

Example 4

A twelve month double-blind placebo-controlled clinical trial is conducted with thirty men and women with renal disease who are undergoing chronic hemodialysis. All patients enter an 8-week control period during which time they receive a maintenance dose of Vitamin D₃ (400 IU/day). After this control period, the patients are randomized into two treatment groups: one group receives a constant dosage of 1α -Vitamin D₂ (u.i.d.; a dosage greater than 3.0 μ g/day) and the other group receives a matching placebo. Both treatment groups receive a maintenance dosage of Vitamin D₃, maintain a normal intake of dietary calcium, and refrain from using calcium supplements. Efficacy is evaluated by pre- and post-treatment comparisons of the two patient groups with regard to (a) direct measurements of intestinal calcium absorption, (b) total body calcium retention, (c) radial and spinal bone mineral density, and (d) determinations of serum calcium and osteocalcin. Safety is evaluated by regular monitoring of serum calcium.

Analysis of the clinical data show that 1α -Vitamin D_2 significantly increases serum osteocalcin levels and intestinal calcium absorption, as determined by direct measurement using a double-isotope technique. Patients treated with this compound show normalized serum calcium levels, stable values for total body calcium, and stable radial and spinal bone densities relative to baseline values. In contrast, patients treated with placebo show frequent hypocalcemia, significant reductions in total body calcium and radial and spinal bone density. An insignificant incidence of hypercalcemia is observed in the treated group.

The foregoing examples demonstrate that 1α -Vitamin D_2 is effective in preventing or restoring the loss of bone mass or bone mineral content while being substantially less toxic than 1α -Vitamin D_3 . It is to be understood that although the foregoing examples detail the use of 1α -Vitamin D_2 , other compounds within the scope of the claims may be readily utilized in the treatment of this invention with essentially equivalent results. For example, 1,25 Vitamin D_2 shows activity equivalent to 1,25 Vitamin D_3 and is also significantly less toxic than its Vitamin D_3 counterpart. Also included within the scope of the claims would be administration of effective dosages of the compound of formula I in conjunction with administration of other Vitamin D compounds, hormones or other agents which have been shown to stimulate bone formation in subjects experiencing or tending toward loss of bone mass or bone mineral content.

Such other Vitamin D compounds would include Vitamin D, 25OH Vitamin D, 1,25 Vitamin D_3 , 1α -Vitamin D_3 and chemical variations which retain the characteristics of these Vitamin D compounds and are contemplated as equivalents.

Such hormones or other agents would include conjugated estrogens or their equivalents, calcitonin, biphosphonates, calcium supplements, cobalamin, pertussis toxin and boron. Possible dose ranges for these co-administered agents are provided in Table 2.

TABLE 2

POSSIBLE ORAL DOSE RANGES FOR VARIOUS AGENTS CO-ADMINISTERED WITH 1α -HYDROXYVITAMIN D_2

AGENT	DOSE RANGES		
	Broad	Preferred	Most Preferred
Conjugated Estrogens or Equivalent (mg/day)	0.3-5.0	0.4 - 2.4	6.6 - 1.2
Sodium Fluoride (mg/day)	5 - 150	30 - 75	40 - 60
Calcitonin (IU/day)	5 - 800	25 - 500	50 - 200
Biphosphonates	50 - 2000	100 - 1500	250 - 1000
Calcium Supplements (mg/day)	250 - 2500	500 - 1500	750 - 1000
Cobalamin (μ g/day)	5 - 200	20 - 100	30 - 50
Pertussis Toxin (mg/day)	0.1 - 2000	10 - 1500	100 - 1000
Boron (mg/day)	0.10 - 3000	1 - 250	2 - 100

Although the examples detail dosage by mouth, it is to be understood that the compounds can be administered in alternative fashions, including intranasally, transdermally, intrarectally, intravaginally, subcutaneously, intravenously, and intramuscularly.

Dosage forms of 1α -Vitamin D_2 can be prepared by dissolving or suspending the compound in, or adsorbing it on, non-toxic pharmaceutically acceptable carriers as is well known in the art. Such carriers may be either solid or liquid such as, for example, corn starch, lactose, sucrose, peanut oil, olive oil, neutral oil, and propylene glycol. If a solid carrier is used, the dosage form of the compound may be tablets, capsules, powders, suppositories, or lozenges. If a liquid carrier is used, soft gelatin capsules, transdermal patches, aerosol sprays, topical creams, syrups or

liquid suspensions, emulsions or solutions may be the dosage form. The dosage forms may also contain excipients, such as preserving, stabilizing, wetting, emulsifying agents, etc.

Bulk quantities of 1α -Vitamin D₂ for the practice of this invention can be readily obtained in accordance with the processes of U.S. Patents Nos. 3,907,843, 4,195,027, 4,202,829, 4,234,495, 4,260,549, 4,555,364, and 4,554,106.

CLAIMS

Having thus described the invention, what we claim is:

1. A method for preventing or reversing loss of bone mass or bone mineral content in mammals displaying or predisposed to developing a bone depletive disorder, comprising: administering to said mammals an effective amount of 1α -Vitamin D_2 .
2. The method according to Claim 1 wherein said mammals are human beings.
3. The method according to Claim 2 wherein said human beings are greater than 45 years of age.
4. The method according to Claim 2 wherein said human beings are women who are hypoestrogenic.
5. The method according to Claim 2 wherein the compound is administered to women during or subsequent to menopause.
6. The method according to Claim 2 wherein said depletive bone disorder is osteoporosis.
7. The method according to Claim 3 or 6 wherein said osteoporosis is post-menopausal osteoporosis.
8. The method according to Claim 3 or 6 wherein said osteoporosis is senile osteoporosis.
9. The method of Claim 2 wherein the compound is administered to women prior to the onset of menopause.
10. The method of Claim 2 wherein 1α -Vitamin D_2 is administered in an amount greater than $3.0 \mu\text{g/day}$.
11. The method of Claim 1 wherein the compound, in solution, in a liquid vehicle ingestible by and non-toxic to said mammals is administered orally in encapsulated form.
12. The method of Claim 1 wherein the compound is administered in combination with at least one hormone or other agent characterized by said hormone's or other agent's ability to reduce loss of bone mass, or bone mineral content in mammals.
13. The method of Claim 12 wherein said hormone or other agent includes other Vitamin D compounds, conjugated

estrogens, sodium fluorides, biphosphonates, cobalamin, pertussin toxin or boron.

14. The method of Claim 1, wherein said administration of said compound is parenteral.
15. The method of claim 14 wherein said administration is by subcutaneous, intramuscular, or intravenous injection, nasopharyngeal or mucosal absorption, or transdermal absorption.
16. The method of claim 1 wherein said administration of said compound is nonparenteral.
17. A pharmaceutical composition comprising the compound 1α -Vitamin D_2 and a pharmaceutically acceptable excipient.
18. A pharmaceutical composition as claimed in Claim 17 which further comprises at least one hormone or other agent's ability to reduce loss of bone mass or bone mineral content in mammals experiencing or tending toward said loss of bone mass or bone mineral content.
19. A pharmaceutical composition as claimed in Claim 18 wherein said hormone or other agent includes other Vitamin D compounds, conjugated estrogens, calcitonin, sodium fluoride, biphosphonates, calcium supplements, cobalamin, pertussin toxin or boron.
20. A method for preventing or restoring in a human, lost bone mass or bone mineral content, said lost bone mass or bone mineral content being related to post-menopausal osteoporosis, which comprises administering to said human:
 - a. 1α -Vitamin D_2 in an amount greater than $3.0 \mu\text{g/day}$; and,
 - b. Limiting the calcium intake of said human to no greater than 800 mg/day .
21. The compound 1α -Vitamin D_2 used as a medicament for the treatment of osteoporosis.
22. A medicine for the treatment of osteoporosis containing as an active ingredient an effective amount of 1α -Vitamin D_2 .

23. A medicine for the treatment of osteoporosis containing as an active ingredient 1α -Vitamin D₂.